Int. Appl. No. : PCT/GB2004/004341
Int. Filing Date : October 13, 2004

AMENDMENTS TO THE CLAIMS

- 1. (Cancelled)
- 2. (Original) A pharmaceutical composition comprising C-peptide together with at least one pharmaceutically acceptable carrier or excipient for administration to a patient as a once daily dose, wherein said once daily dose does not include a continuous administration or the presence of release rate-controlling agents for the treatment of diabetes and/or microvascular diabetic complications.
- 3. (Currently amended) Method A method of treating diabetes and/or microvascular diabetic complications comprising administering C-peptide or a pharmaceutical composition comprising C-peptide to a patient in a once daily dose, wherein said once daily dose does not include a continuous administration or the presence of release rate-controlling agents.
- 4. (Currently amended) Use, The pharmaceutical composition or method according to Claim 2 any one of claims 1 to 3 wherein the C-peptide is human C-peptide
- 5. (Currently amended) Use, The pharmaceutical composition or method according to Claim 2 any one of claims 1 to 4 wherein said C-peptide is a the fragment EGSLQ (SEQ ID NO:[[-]] 2).
- 6. (Currently amended) Use, The pharmaceutical composition or method according to Claim 2 any one of claims 1 to 5 wherein the patient is a human.
- 7. (Currently amended) Use, The pharmaceutical composition or method according to Claim 2 any one of claims 1 to 6 wherein the medicament contains 100 to 1800 nmol of C-peptide.
- 8. (Currently amended) Use, The pharmaceutical composition or method according to Claim 2 any one of claims 1 to 7 wherein the medicament is an uncompromised aqueous solution.
- 9. (Currently amended) Use, The pharmaceutical composition or method according to Claim 2 any one of claims 1 to 8 wherein said complications are diabetic nephropathy, retinopathy or neuropathy.
- 10. (New) The method according to Claim 3, wherein the C-peptide is human C-peptide

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11. (New) The method according to Claim 3, wherein said C-peptide is a fragment EGSLQ (SEQ ID NO: 2).

- 12. (New) The method according to Claim 3, wherein the patient is a human.
- 13. (New) The method according to Claim 3, wherein the medicament contains 100 to 1800 nmol of C- peptide.
- 14. (New) The method according to Claim 3, wherein the medicament is an uncompromised aqueous solution.
- 15. **(New)** The method according to Claim 3, wherein said complications are diabetic nephropathy, retinopathy or neuropathy.